PATENT APPLN. NO. 10/534,874 SUBMISSION UNDER 37 C.F.R. §1.114 PATENT NON-FINAL

## IN THE CLAIMS:

- 1. (currently amended) A liposome to which a polyalkylene glycol and wild type human serum albumin are bonded selected from the group consisting of:

  (a) a liposome to which each of a polyalkylene glycol and a non-modified serum albumin is bonded, wherein the non-modified serum albumin is bonded to the liposome via a reactive intervening group;

  (b) a liposome to which a serum albumin is bonded via a polyalkylene glycol, wherein the serum albumin is bonded to the polyalkylene glycol via a reactive intervening group; and

  (c) a liposome wherein the liposome and a polyalkylene glycol are bonded to a serum albumin via reactive intervening groups at a different site.
- 2. (original) The liposome according to claim 1, wherein a physiologically active ingredient is further contained.
- 3. (original) The liposome according to claim 2, wherein the physiologically active ingredient is a pharmaceutically active ingredient.

- 4. (original) The liposome according to claim 3, wherein the pharmaceutically active ingredient is an antitumor agent.
- 5. (currently amended) A pharmaceutical composition containing the liposome mentioned recited in claim 2.
- 6. (previously presented) The pharmaceutical composition according to claim 5, wherein the composition is in an injection form.
- 7. (currently amended) A method for treatment of cancer, which comprises administering a pharmaceutical composition comprising a liposome to which a polyalkylene glycol and wild type human serum albumin are bonded and in which an antitumor agent is contained.
- 8. (currently amended) A method of extending the in vivo retention time of a physiologically active ingredient contained in a liposome comprising binding the liposome to a polyalkylene glycol and wild type human serum albumin.

- 9. (currently amended) A process for the production of the liposome comprising a step selected from the group consisting of (A), (B), (C), (D), (E) and (F):[[,]]
- (A) a step of bonding a liposome having containing a compound represented by the following formula (1):

(wherein R is an acyl group derived from a fatty acid having 2 to 35 carbon atoms) and 1,2-distearol-sn-glycero-3-phosphoethanolamine bonded to a polyalkylene glycol (PEG-DSPE) as constituent lipids to wild type human serum albumin;

(B) a step of bonding a liposome having containing a compound represented by the following formula (2):

$$\begin{array}{c|c}
CH_2 \longrightarrow OR \\
RO \longrightarrow CH & O \\
CH_2 \longrightarrow O \longrightarrow P \longrightarrow OCH_2CIL_NHCOCH_2CH_2 \longrightarrow S \longrightarrow S
\end{array}$$
(2)

(wherein R has the same meaning as defined above) and 1,2-distearol-sn-glycero-3-phosphoethanolamine bonded to a polyalkylene glycol (PEG-DSPE) as constituent lipids to a compound represented by the formula (3):

$$(Alb-NH)-CO-CH2-CH2-SH$$
 (3)

(wherein Alb-NH is a group formed by removing one hydrogen atom of the amino group from [[an]] a wild type human serum albumin molecule represented by Alb-NH<sub>2</sub>);

(C) a step of bonding a liposome having containing a compound represented by the following formula (4):

(4)

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(wherein n is an integer of 5 to 100,000 and R has the same meaning as defined above) as a constituent lipid is bonded to a compound represented by the formula (5):

$$(Alb-NH)-CO-CH2-SH (5)$$

(wherein Alb-NH has the same meaning as defined above);

(D) a step of bonding incorporating a compound represented by the following formula (6):

$$\begin{array}{c} CH_2 \longrightarrow OR \\ RO \longrightarrow CH \\ CH_2 \longrightarrow O \longrightarrow P \longrightarrow OCH_2CH_2NHCO \\ OH \\ OCH_2 \end{array} CH_2 \longrightarrow CH_2CO \longrightarrow (NH-Alb) \\ OCH_2 CH_2 \longrightarrow OCH_2CH_2 \longrightarrow OCH_2 \longrightarrow OCH_2CH_2 \longrightarrow OCH_2 \longrightarrow OCH_2$$

(wherein n, R and Alb-NH have each the same meaning as defined above) into a liposome;

(E) a step of bonding a liposome having containing the compound represented by the above formula (1) as a constituent lipid to a compound represented by the following formula (7):

$$S - CH_2CO - (NH-Alb-NIL)$$
 $CH_3OCH_2CH_2 - OCH_2CH_2 - OCH_2CH_$ 

(wherein  $-NH-Alb-NH_2$  is a group formed by removing one hydrogen atom from one of the amino groups of an albumin molecule represented by  $H_2N-Alb-NH_2$ , and n has the same meaning as defined above); or

(F) a step of bonding a liposome having containing the compound represented by the above formula (2) as a constituent lipid to a compound represented by the following formula (8):

$$CH_{2}CO \longrightarrow (NH-Alb-NH) \longrightarrow COCH_{2}CH_{2}SH$$

$$CH_{3}OCH_{2}CH_{2} \longrightarrow CH_{2}CH_{2} \longrightarrow N$$

$$O$$

$$(8)$$

(wherein -NH-Alb-NH- is a group formed by removing one hydrogen atom from each of the two amino groups of an albumin molecule represented by the formula  $H_2N-Alb-NH_2$ , and n has the same meaning as defined above).

- 10. (previously presented) A pharmaceutical composition containing the liposome mentioned in claim 3.
- 11. (previously presented) A pharmaceutical composition containing the liposome mentioned in claim 4.
- 12. (previously presented) The pharmaceutical composition according to claim 10, wherein the composition is in an injection form.
- 13. (previously presented) The pharmaceutical composition according to claim 11, wherein the composition is in an injection form.
  - 14. 16. (canceled)